Pioneering the future of biomanufacturing

ENTRI
ENABLING TRANSLATION

ENTRI supports companies seeking flexible, ready-to-use cGMP cleanroom space, including wet labs and office space, with options for partial or full facility use. Whether you are a biotech, pharma or CDMO looking for space, ENTRI is for you.

As a purpose-built, stand-alone cGMP manufacturing facility, ENTRI provides scalable manufacturing infrastructure, cGMP capability and technical support for a range of products, including biologics (such as ADCs and mAbs), cell and gene therapies (such as CAR T-cell therapies), RNA-based products (such as mRNA-LNPs) and drug-device combination products, with additional modalities supported subject to facility fit.

Supporting clinical and commercial manufacturing needs

Supporting you from technology transfer through to clinical supply, ENTRI enables you to undertake analytical and process development, drug substance and potentially drug product manufacturing, on-site QC testing, with the capability for scalable commercial manufacturing.

Flexible support to establish, scale and execute

ENTRI's technical team can support areas including equipment procurement and qualification, quality and regulatory activities, person-in-plant, and workforce development (subject to requirements).

Based on technical alignment, this can be complemented by a more traditional CDMO-style support as you build your local workforce and scale your operations.

Our technical team brings extensive industry expertise across cGMP manufacturing, operations, validation, and quality, with an average of more than 20 years' experience working with leading CDMOs and global pharma.

A Strategic APAC Base with Secure Capacity, Control and Global Reach

ENTRI
ENABLING TRANSLATION

ENTRI offers a unique blend of in-house control and external site benefits:

- **Enhanced Control:** Maintain full oversight of your critical production processes and manufacturing intellectual property (IP).
- **Flexibility:** Easily adapt to changing production demands, schedules, and timelines.
- **Dedicated Capacity:** Secure guaranteed manufacturing capacity to use across your pipeline assets.
- **Compliance:** Benefit from facilities designed and built to meet cGMP and global regulatory standards.
- **Financial Benefits:** Avoid costly investments to build new facilities or upgrade existing buildings.
- **Strategic APAC Location:** Leverage Australia's strong, stable supply chains and trade ties with key international markets, including the US and Europe.

Benefits of working in Australia

Located in Brisbane, Australia, ENTRI is part of the Translational Research Institute ecosystem within the Boggo Road Innovation Junction (BRIJ), a growing biomedical innovation cluster that connects manufacturing, research, clinical, and commercial capabilities to enable clinical translation and product commercialisation.

Companies can access Australia's R&D Tax Incentive, potentially offering up to a 48.5% cash refund on manufacturing and R&D costs for eligible companies.

Australia offers one of the fastest regulatory paths to initiate clinical trials globally, without requiring an IND.

Find out more: entri.com.au

Proudly funded by the Queensland Government and TRI.

Queensland Government